



Form PTO-1449 (modified)		Atty. Docket No. GOUD:023USD2	Serial No. 10/664,423
Office of Patents and Publications for Applicant's		Applicant Guy A. Rouleau <i>et al.</i>	
INFORMATION DISCLOSURE STATEMENT (Use several sheets if necessary)		Filing Date: September 17, 2003	Group: 1649 Unknown
U.S. Patent Documents <i>See Page 1</i>	Foreign Patent Documents <i>See Page 1</i>	Other Art <i>See Page 1</i>	

U.S. Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Name	Class	Sub Class	Filing Date of App.

Foreign Patent Documents

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Exam. Init.	Ref. Des.	Citation
DK	C73	Ahmed et al., "Primary structure, chromosomal localization, and functional expression of a voltage-gated sodium channel from human brain," Proc. Natl. Acad. Sci. USA, 89:8220-8224, 1992.
DK	C74	Lu and Brown, "Isolation of a human-brain sodium-channel gene encoding two isoforms of the subtype III α -subunit," J. Mol. Neuro., 10:67-70, 1998.

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DK	A1	5,223,409	6-29-93	Ladner et al.	435	69.7	3-1-91

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	C5	Barker et al., "GABA actions on the excitability of cultured CNS neurons," <i>Neurosci. Lett.</i> , 47:313-318, 1984.
	C6	Bar-Sagi et al., "Negative modulation of sodium channels in cultured chick muscle cells by the channel activator batrachotoxin," <i>J. Biol. Chem.</i> , 260:4740-4744, 1985.
	C7	Baulac et al., "A second locus for familial generalized epilepsy with febrile seizures plus maps to chromosome 2q21-q33," <i>Am. J. Hum. Genet.</i> , 65:1078-1085, 1999.
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	C9	Berkovic et al., "Epilepsies in twins: genetics of the major epilepsy syndromes," <i>Ann. Neurol.</i> , 43:435-445, 1998.
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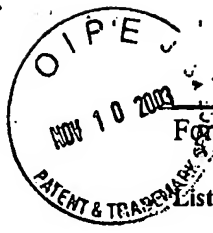
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	C13	Charlier et al., "A pore mutation in a novel KQT-like potassium channel gene in an idiopathic epilepsy family," <i>Nat. Genet.</i> , 18:53-55, 1998.
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	C16	Cho et al., "An Unnatural Biopolymer," <i>Science</i> , 261:1303-1305, 1993.
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	C18	Corey et al., "The occurrence of epilepsy and febrile seizures in Virginian and Norwegian twins," <i>Neurology</i> , 41:1433-1436, 1991.
	C19	Cull et al., "Screening for receptor ligands using large libraries of peptides linked to the C terminus of the lac repressor," <i>Proc. Natl. Acad. Sci. USA</i> , 89:1865-1869, 1992.
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	C21	DeWitt et al., "'Diversomers': an approach to nonpeptide, nonoligomeric chemical diversity," <i>Proc. Natl. Acad. Sci. USA</i> , 90:6909-6913, 1993.
	C22	Elliot et al., "Bin1 functionally interacts with Myc and inhibits cell proliferation via multiple mechanisms," <i>Oncogene</i> , 18:3564-3573, 1999.
	C23	Elmslie et al., "Genetic mapping of a major susceptibility locus for juvenile myoclonic epilepsy on chromosome 15q," <i>Hum. Mol. Genet.</i> , 6:1329-1334, 1997.
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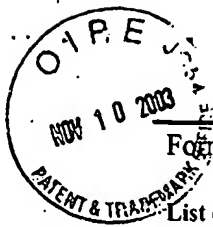
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	C26	Escayg et al., "22," <i>Nat. Genet.</i> , 24:343-345, 2000. Mutations of SCN1A, encoding a neuronal sodium channel, in two families with GEFS+
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	C62	Scott et al., "Searching for peptide ligands with an epitope library," <i>Science</i> , 249:386-390, 1990.
	C63	Sillampää et al., "Genetic factors in epileptic seizures: evidence from a large twin population," <i>Acta Neurol. Scand.</i> , 84:523, 1991.
	C64	Singh et al., "A novel potassium channel gene, KCNQ2, is mutated in an inherited epilepsy of newborns," <i>Nat. Genet.</i> , 18:25-29, 1998.
✓	C65	Sjolander et al., "Integrated fluid handling system for biomolecular interaction analysis," <i>Anal. Chem.</i> , 63:2338-2345, 1991.

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DK	C66	Steinlein et al., "A missense mutation in the neuronal nicotinic acetylcholine receptor alpha 4 subunit is associated with autosomal dominant nocturnal frontal lobe epilepsy," <i>Nat. Genet.</i> , 11:201-203, 1995.
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	C68	Tamaskovic et al., "Enzyme-linked immunosorbent assay for the measurement of JNK activity in cell extracts," <i>Biological Chemistry</i> , 380:569-578, 1999.
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	C70	Wallace et al., "Febrile seizures and generalized epilepsy associated with a mutation in the Na ⁺ -channel beta1 subunit gene SCN1B," <i>Nature Genet.</i> , 19:366-370, 1998.
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